DE LA CRUZ CAMPHOR - camphor ointment
DLC Laboratories, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

CAMPHOR Ointment 11%

Active Ingredient

Camphor, USP 11%

Purpose

External analgesic

Uses

for the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises and sprains

Warnings

For external use only

When using this product

avoid contact with eyes or mucous membranes

do not apply to wounds or damaged skin

do not bandage tightly or use with a heating pad

Stop use and consult a doctor if

condition worsens or if symptoms persist for more than 7 days

symptoms clear up and occur again within a few days.

excessive skin irritation develops

If pregnant or breastfeeding,

consult a doctor before use.

KEEP OUT OF THE REACH OF CHILDREN.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

adults and children 2 years of age or older, apply to affected area not more than 3 to 4 times daily

children under 2 years of age: consult a doctor

Inactive ingredient

polyethylene glycol

Questions?

1-800-858-3889 or www.dlclabs.com

De La Cruz

CAMPHOR
Ointment 11%

Pain relieving rub

2.5 OZ (70.9g)

FAST, PENETRATING RELIEF FOR:

Muscle and joint pain

Backaches and arthritis

Strains and sprains

Itching

NON-IRRITATING

WATER WASHABLE

NO PARABENS OR ARTIFICIAL FRAGRANCES OR COLORS

Manufactured by:

De La Cruz Products

A Division of DLC Laboratories, Inc.

Paramount, CA 90723 USA

Questions: 1-800-858-3889

www.dlclabs.com (c) DLC

PRINCIPAL DISPLAY PANEL - 70.9 g Jar Label

De La Cruz®

Clinically Tested*

CAMPHOR
OINTMENT 11%

Pain Relieving Rub

2.5 OZ (70.9 g)

Datos Medicinales

Ingrediente activo — *Propósito*
Alcanfor, USP 11%..... Analgésico externo

Usos para el alivio temporal de leves dolores musculares y de articulaciones asociados con: ■ simples dolores de espalda ■ artritis ■ torceduras ■ esguinces ■ moretones

Advertencias
Para uso externo únicamente

Cuando use este producto
■ evite el contacto con los ojos o las membranas mucosas
■ no aplique sobre heridas o piel lastimada
■ no vende ajustadamente ni use junto a una almohadilla térmica

Deje de usar y consulte con un médico si
■ la condición empeora o si los síntomas persisten por más de 7 días ■ los síntomas desaparecen y aparecen nuevamente a los pocos días ■ se desarrollara una excesiva irritación en la piel

Si está embarazada o lactando, consulte con un médico antes de usar.

MANTÉNGASE FUERA DEL ALCANCE DE LOS NIÑOS. En caso de ser ingerido, pida ayuda médica o llame inmediatamente a un Centro de Control de Envenenamientos.

Instrucciones
■ adultos y niños mayores de 2 años de edad, aplique al área afectada no más de 3 a 4 veces al día ■ niños menores de 2 años de edad: consulte con un médico

Ingrediente inactivo
glicol polietileno

¿Preguntas?
1-800-858-3889 o www.dlclabs.com

Drug Facts

Active ingredient — *Purpose*
Camphor, USP 11%..... External analgesic

Uses for the temporary relief of minor aches and pains of muscles and joints associated with: ■ simple backache ■ arthritis ■ strains ■ sprains ■ bruises

Warnings
For external use only

When using this product
■ avoid contact with eyes or mucous membranes ■ do not apply to wounds or damaged skin ■ do not bandage tightly or use with a heating pad

Stop use and consult a doctor if
■ condition worsens or if symptoms persist for more than 7 days ■ symptoms clear up and occur again within a few days ■ excessive skin irritation develops

If pregnant or breast-feeding, consult a doctor before use.

KEEP OUT OF THE REACH OF CHILDREN. If swallowed, get medical help or contact a Poison Control Center right away.

Directions
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Inactive ingredient
polyethylene glycol

Questions?
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DE LA CRUZ CAMPHOR

camphor ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24286-1521
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	11 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24286-1521-2	70.9 g in 1 JAR; Type 0: Not a Combination Product	07/26/2012	
2	NDC:24286-1521-5	155.9 g in 1 JAR; Type 0: Not a Combination Product	03/10/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/26/2012	

Labeler - DLC Laboratories, Inc. (093351930)

Establishment

Name	Address	ID/FEI	Business Operations
DLC Laboratories, Inc.		093351930	MANUFACTURE(24286-1521) , LABEL(24286-1521)

Revised: 3/2021 DLC Laboratories, Inc.